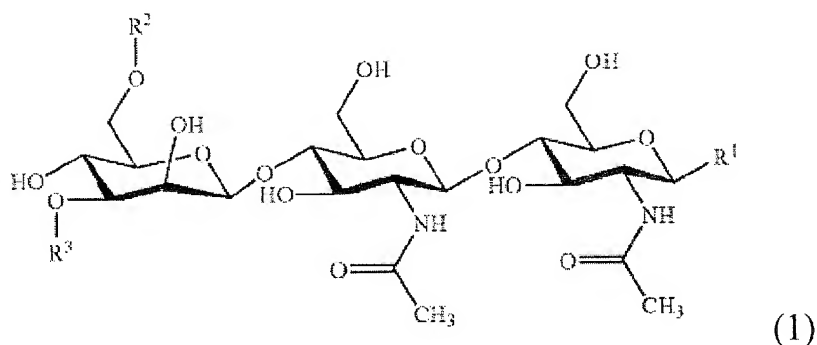


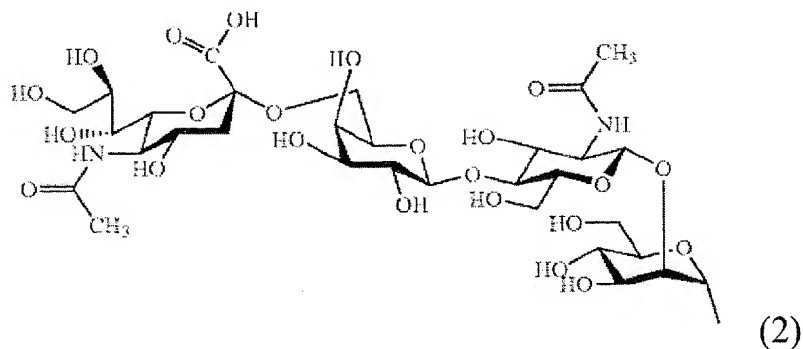
IN THE CLAIMS:

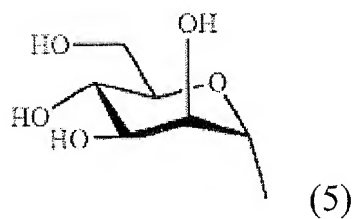
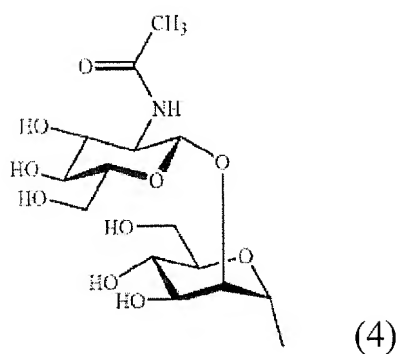
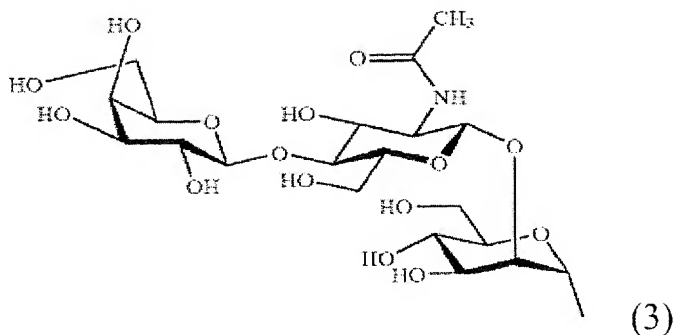
1-3. (Canceled).

4. (Currently amended) A glycopeptide ~~resistant to sugar hydrolase which cleaves the reducing terminal of an oligosaccharide from a peptide~~ comprising an aminated complex-type oligosaccharide of the formula (1)



wherein R^1 is ~~$\text{-NH-(CO)-CH}_2\text{X}$, $\text{-NH-(CO)-(CH}_2\text{)}_b\text{-CH}_2\text{X}$, isothiocyanate group, $\text{-NH-(CO)}_a\text{-(CH}_2\text{)}_b\text{-CO}_2\text{H}$ or $\text{-NH-(CO)}_a\text{-(CH}_2\text{)}_b\text{-CHO}$~~ , X being a halogen atom, ~~a being 0 or 1, b being an integer of 1 to 4~~, R^2 and R^3 are a hydrogen atom or a group of the formulae (2) to (5) and may be the same or different, except that R^2 and R^3 are not both hydrogen or the formula (5) at the same time and when one of R^2 and R^3 is hydrogen, the other is not the formula (5),





wherein the glycopeptide has about 12 times higher resistance to Peptide-N Glycosidase F (PNGase F) than a glycopeptide comprising an asparagine-linked oligosaccharide, and the aminated complex-type oligosaccharide binds to a thiol group of a peptide by displacement of halogen X of NH-(CO)-CH₂X and a thiol group of a peptide bonded thereto.

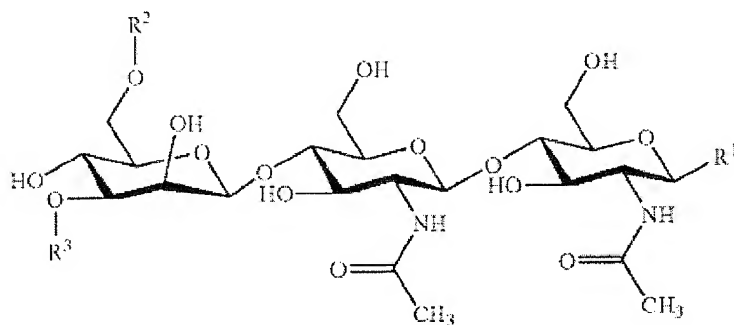
5. (Canceled).

6. (Original) A glycopeptide as defined in claim 4 wherein the glycopeptide is an antibody.

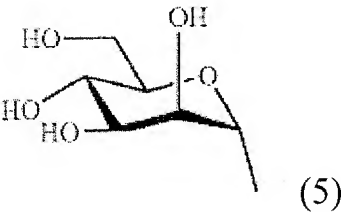
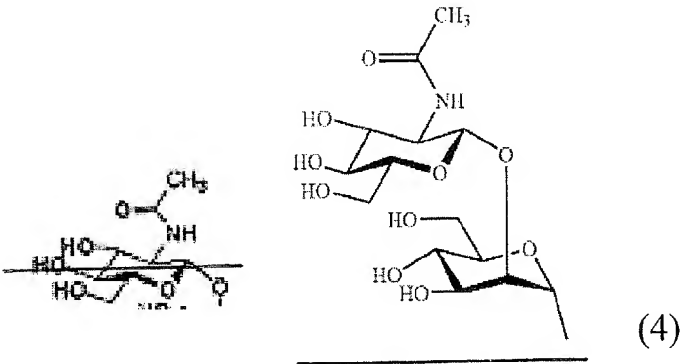
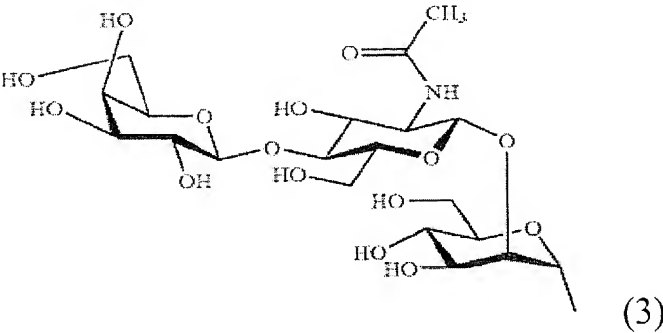
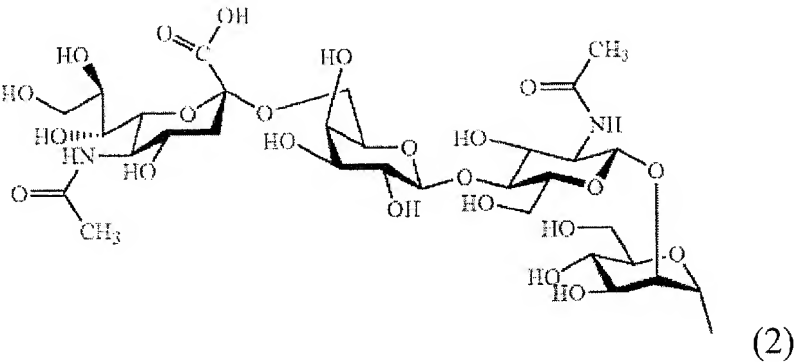
7. (Currently amended) A process for preparing a uniform glycopeptide composition comprising steps of (a) and (b) that are performed at the same time,

(a) cleaving an asparagine-linked oligosaccharide ~~from a glycopeptide from a peptide by Peptide-N Glycosidase F (PNGase F) sugar hydrolase which cleaves the reducing terminal of an oligosaccharide from a peptide, wherein the resulting peptide has a thiol group,~~
 and

(b) bonding an aminated complex-type oligosaccharide of the formula (1)



wherein R^1 is ~~$-\text{NH}-(\text{CO})-\text{CH}_2\text{X}$, $-\text{NH}-(\text{CO})-(\text{CH}_2)_b-\text{CH}_2\text{X}$, isothiocyanate group, $-\text{NH}-(\text{CO})_a-(\text{CH}_2)_b-\text{CO}_2\text{H}$ or $-\text{NH}-(\text{CO})_a-(\text{CH}_2)_b-\text{CHO}$, X being a halogen atom, a being 0 or 1, b being an integer of 1 to 4, R^2 and R^3 are a hydrogen atom or a group of the formulae (2) to (5) and may be the same or different, except that R^2 and R^3 are not both hydrogen or the formula (5) at the same time and when one of R^2 and R^3 is hydrogen, the other is not the formula (5),~~



to the thiol group of the resulting peptide by displacement of halogen X of -NH-(CO)-CH₂X.

8. (Previously presented) A glycopeptide prepared according to the process of claim 7, the glycopeptide prepared being an antibody.